

510(k) Summary

NOV 17 2005

K053042

OFFICIAL CONTACT: Lisa A. Ewing
Regulatory Affairs Specialist
MEDRAD, INC.
One Medrad Drive
Indianola, PA 15051
(412) 767-2400 Ext. 3780

CLASSIFICATION NAME: Magnetic Resonance Diagnostic Device
(21 CFR 892.1000, Product Code MOS)

COMMON NAME(S): Magnetic Resonance Diagnostic Accessory

PROPRIETARY NAME: MEDRAD 1.5T Pelvic Imaging System Interface Device

PREDICATE DEVICE: MEDRAD, INC. 1.5T ATD-T Interface Device
(K974438)

INTENDED USE: The 1.5T Pelvic Imaging System is a receive-only coil intended for use as a Magnetic Resonance Diagnostic Device (MRDD) for high-resolution magnetic resonance imaging, including spectroscopy, of the human colon, cervix or prostate and surrounding pelvic tissue. The purpose of the interface device for the 1.5T Pelvic Imaging System is to provide interface and support functions to allow the use of MEDRAD 1.5T disposable endorectal coils with GE Signa EXCITE 1.5T MRI scanner systems and body array coils. Only trained healthcare professionals are intended to operate this device.

DEVICE DESCRIPTION AND COMPARISON TO UNMODIFIED PREDICATE:

The MEDRAD 1.5T Pelvic Imaging System Interface Device maintains a similar intended use, similar operational parameters, similar labeling and is used in a manner similar to the predicate device.

The following comparison table identifies the similarities and differences between the new device and the predicate device.

Comparison of Features and Principles of Operation in MEDRAD 1.5T ATD-T (Predicate) and MEDRAD 1.5T Pelvic Imaging System Interface Device (Proposed)

Feature	(Predicate) MEDRAD 1.5T ATD-T	(Proposed) MEDRAD Pelvic Imaging System Interface Device
Coil Type	Receive-only surface coil.	Receive-only surface coil.
Scanner Interface	1.5T GE Signa scanners	1.5T GE Signa EXCITE scanners
Coil Interfaces	MEDRAD 1.5T Endorectal Coils, GE 4-channel Torso Array	MEDRAD 1.5T Endorectal Coils, GE 8- or 12-channel Body Array
Decoupling	Active and passive	Active and passive
RF Signal Pre-Amplification	Performed by the scanner for both the Endorectal Coil and the Torso Array.	Performed by the interface device for the Endorectal Coil (RF output from Body Array is already preamplified).
Tuning	Requires tuning to be set in production since ATD-T uses system preamplifier for signal amplification.	Does not require tuning to be set in production since interface device has a preamplifier.



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Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

Ms. Lisa A. Ewing
Regulatory Affairs Specialist
MEDRAD, Inc.
One Medrad Drive
INDIANOLA PA 15051-0780

Re: K053042
Trade/Device Name: MEDRAD 1.5T Pelvic Imaging
System Interface Device
Regulation Number: 21 CFR 892.1000
Regulation Name: Magnetic resonance
diagnostic device
Regulatory Class: II
Product Code: MOS
Dated: October 27, 2005
Received: October 28, 2005

Dear Ms. Ewing:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Office of Compliance at one of the following numbers, based on the regulation number at the top of this letter:

21 CFR 876.xxxx	(Gastroenterology/Renal/Urology)	240-276-0115
21 CFR 884.xxxx	(Obstetrics/Gynecology)	240-276-0115
21 CFR 892.xxxx	(Radiology)	240-276-0120
Other		240-276-0100

Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR 807.97). You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its Internet address <http://www.fda.gov/cdrh/industry/support/index.html>.

Sincerely yours,



Nancy C. Brogdon
Director, Division of Reproductive,
Abdominal, and Radiological Devices
Office of Device Evaluation
Center for Devices and Radiological Health

Enclosure

Indications for Use

510(k) Number: K 053042

Device Name: MEDRAD 1.5T Pelvic Imaging System Interface Device

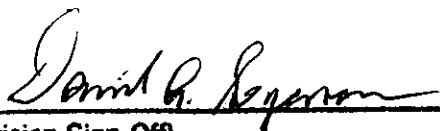
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Prescription Use X ~~AND/OR~~ Over-The-Counter Use _____
(Part 21 CFR 801 Subpart D) (21 CFR 801 Subpart C)

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE OF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)


(Division Sign-Off)
Division of Reproductive, Abdominal,
and Radiological Devices
510(k) Number K 053042

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